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(54) Title: IN VITRO ANTIBODY AFFINITY MATURATION USING ALANINE SCANNING MUTAGENESIS			
(57) Abstract <p>A method of mutagenizing antibodies to produce modified antibodies, modified antibodies, DNA encoding the modified antibodies as well as diagnostic kits and pharmaceutical compositions comprising the antibodies or DNA are provided.</p>			

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TITLE OF THE INVENTION

IN VITRO ANTIBODY AFFINITY MATURATION USING ALANINE SCANNING MUTAGENESIS

CROSS-RELATED TO OTHER APPLICATIONS

5 This is a continuation of U.S. Serial No. 08/206,076 filed March 4, 1994, now pending.

BRIEF DESCRIPTION OF INVENTION

10 A method of mutagenizing antibodies to produce modified antibodies, modified antibodies, DNA encoding the modified antibodies as well as diagnostic kits and pharmaceutical compositions comprising the antibodies or DNA are provided. The method of the invention is a systematic means to achieve *in vitro* antibody maturation and uses alanine scanning mutagenesis. The 15 invention is particularly exemplified with a set of single chain Fv (scFv) antibodies obtained by this technique. The resulting antibodies are directed against the V3 loop of HIV gp120, and show altered off-rates against the antigen compared to the starting antibody. Of particular interest are the altered antibodies which 20 show improved (slower) off-rates to the antigen. Observed improvements have been as high as eleven-fold over wild-type.

SUMMARY OF THE INVENTION

25 A method of mutagenizing antibodies to produce modified antibodies, modified antibodies, DNA encoding the modified antibodies as well as diagnostic kits and pharmaceutical compositions comprising the antibodies or DNA are provided.

BRIEF DESCRIPTION OF THE DRAWINGS

30 Figure 1. Alanine-Scanning Mutagenesis. Each of the 27 amino acids in VH CDR3 of scFv P5Q was converted to alanine by site-directed mutagenesis. *E. coli* clones were induced to express scFv with IPTG. Single chain Fv, which is targeted to the periplasmic space by the fd phage gene3 signal sequence, was

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extracted with EDTA. Periplasmic extracts were analyzed by BIACore™, which measures antibody-antigen affinity by surface plasmon resonance (Fägerstam, 1991), and off-rates determined against an HIV gp120 V3 loop peptide. Results of the alanine scan, relative to P5Q, fall into four classes: i) slower off-rate, ii) faster off-rate, iii) no binding, and iv) minor or no change in off-rate. Standard deviation is  $\pm 25\%$ .

Figure 2. Amino Acid Randomization: Position 107. Arginine at position 107 was mutated to all amino acids by site-directed mutagenesis. Single chain Fv extracts were analyzed by BIACore. Percent change in off-rates is shown relative to P5Q.

Figure 3. Amino Acid Randomization: Position 111. Glutamic acid at position 111 was mutated to all amino acids by site-directed mutagenesis. Single chain Fv extracts were analyzed by BIACore. Percent change in off-rates is shown relative to P5Q.

Figure 4. Amino Acid Randomization: Position 112. Aspartic acid at position 112 was mutated to all amino acids by site-directed mutagenesis. Single chain Fv extracts were analyzed by BIACore. Percent change in off-rates is shown relative to P5Q.

Figure 5. Additive Effect of Combining Optimized Residues. A double mutant, containing the optimized residues, was constructed and analyzed by BIACore. Percent change in off-rates is shown relative to P5Q.

Figure 6. Nucleotide and amino acid sequences of scFv P5Q with c-myc tail.

#### DETAILED DESCRIPTION OF THE INVENTION

The gp120 V3 domain of human immunodeficiency virus-1 (HIV-1) is a disulfide-linked closed loop of approximately 30 amino acids. The loop, in either native or synthetic form, binds to and elicits anti-HIV-1 antibodies.

The present invention relates to modified antibodies and methods of making modified. The invention is exemplified with modified HIV-1 immunoglobulins and methods of making these

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modified HIV-1 immunoglobulins. The modified immunoglobulins of the present invention contain an altered complementary determining region 3 (CDR3) of HIV-1 neutralizing antibody.

5 The present invention also comprises a method of preventing infection through the administration of a modified antibody to a suitable host. In one embodiment of the invention, the treatment or prevention of HIV infection through the administration of the modified HIV-1 immunoglobulin is described.

10 The present invention also comprises diagnostic kits useful for the detection or characterization of an antigen. Reagents for the kits may include DNA molecules encoding the modified antibodies or the modified antibodies or combinations thereof.

15 A method of mutagenizing antibodies to produce modified antibodies, modified antibodies, DNA encoding the modified antibodies as well as diagnostic kits and pharmaceutical compositions comprising the antibodies or DNA are provided. The method of the invention is a systematic means to achieve *in vitro* antibody maturation and uses alanine scanning mutagenesis. The invention is particularly exemplified with a set of single chain Fv (scFv) antibodies obtained by this technique. The resulting antibodies are directed against the V3 loop of HIV gp120, and show altered off-rates against the antigen compared to the starting antibody. Of particular interest are the altered antibodies which show improved (slower) off-rates to the antigen. Observed improvements have been as high as eleven-fold over wild-type.

20 25 Maturation was achieved through an alanine scan of complementary determining region 3 (CDR3) to identify positions critical to antigen binding. Critical positions were then randomized to identify amino acids that provided the slowest off-rates. Finally, clones were optimized through the combining of mutations.

30 The underlying principle of the method is the physical and chemical neutrality of alanine. Alanine is substituted throughout a stretch of amino acids, and its effects on binding (such as off-rate and on-rate) are evaluated using conventional methods. The number

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of positions likely to be identified in this manner is relatively small. Once identified, these key positions may be randomized to all amino acids to identify the best amino acid solution at the position. Because all manipulations and evaluations are conducted *in vitro*,  
5 physiological bias is limited.

Present methods of *in vitro* antibody maturation are essentially random procedures in which the researcher generates clones with amino acid substitutions and evaluates them. The problem is that the number of substitutions necessary for a thorough  
10 evaluation is extremely large. For example, if one were to evaluate all random substitutions in CDR3, a region typically twenty-five residues in length, one would have to examine  $9 \cdot 10^{27}$  possibilities. This is beyond the capabilities of present technologies.

Alanine scanning maturation enables the rapid  
15 identification of residues most likely to be important in binding. Using the example of a twenty-five residue stretch cited above, only twenty-five substitutions would be necessary. From this initial screen, amino acid positions likely to be critical to binding may be identified. The critical residues may then be randomized to identify  
20 the amino acids that optimize binding. Using this method, scFv antibodies with dissociation rates greater than ten-fold slower than the original scFv have been created.

Previous work in *in vitro* antibody maturation used one of two general approaches. In one approach, PCR recombination is  
25 used to substitute all or part of the VH and VL genes into libraries of scFv clones. In the second approach, random mutations are made throughout a CDR region of a scFv clone by the use of degenerate oligonucleotides. In both cases, clones were expressed as a phage fd gene 3 fusion surface protein. Higher affinity clones were identified  
30 using a panning assay followed by clonal purification of the phage.

Each approach has drawbacks. The PCR method is cumbersome, limited to the sequences of the B cell population, is essentially random in nature, and may introduce unwanted mutations through the PCR recombination step. The randomization approach

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produces only a small fraction of the possible CDR changes. Neither approach allows immediate determination of changes in binding affinity because it is necessary to first generate an enriched population of suitable clones through panning. Both approaches detect only changes which result in improved binding; they do not identify positions for which the change weakened the binding. The latter class of change may include critical binding residues in which the appropriate amino acid solutions leads to improvement.

The method disclosed herein is systematic, thorough and unlikely to introduce unexpected or undesired mutations. All manipulations are done *in vitro*, which minimizes bias due to selection steps. Evaluation of clones is quantitative. In some cases, a key amino acid position may display poorer binding with alanine, but subsequent randomization may yield an amino acid solution which enables improved binding. Such mutations would not be detected by previous methods. Because the method of the present invention does not require phage expression for panning, the method can be used on scFVs, Fabs, and full length antibodies. Use is not restricted to a scFv for phage expression. Using the approach of the present invention, an anti-HIV V3 loop antibody was improved approximately eleven-fold.

Alanine scanning maturation of antibodies is a general method which may be used to improve binding of antibodies to their cognate antigens. The method has been used to identify critical residues in the scFv 447 which can be introduced into MAAb447. Such changes may lead to significant improvement of the binding affinity of MAAb447 against multiple species of HIV gp120 isolates. This improvement may increase the neutralization capability of the antibody, and significantly lower the effective dose.

Although the method and antibodies of the present invention are exemplified with scFv antibodies, it is readily apparent to those skilled in the art that the method may be used with other types of antibodies or with antibodies targetted against different

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epitopes or antigens. Other types of antibodies include but are not limited to fragments of antibodies and full-length antibodies.

5 The molecular biology and immunological techniques of the present invention can be performed by standard techniques well-known in the art. See, for example, in Maniatis, T., Fritsch, E.F., Sambrook, J., *Molecular Cloning: A Laboratory Manual* (Cold Spring Harbor Laboratory, Cold Spring Harbor, New York, 1982).

10 Because the genetic code is degenerate, more than one codon may be used to encode a particular amino acid, and therefore, the amino acid sequence can be encoded by any of a set of similar DNA oligonucleotides.

15 The cloned DNA molecules obtained may be expressed by cloning the gene encoding the altered antibody into an expression vector containing a suitable promoter and other appropriate transcription regulatory elements, and transferred into prokaryotic or eukaryotic host cells to produce recombinant modified antibodies. Techniques for such manipulations are well-known in the art.

20 In order to simplify the following Examples and the Detailed Description, certain terms will be defined.

25 Expression vectors are defined herein as DNA sequences that are required for the transcription of cloned copies of genes and the translation of their mRNAs in an appropriate host. Such vectors can be used to express eukaryotic genes in a variety of hosts such as bacteria, bluegreen algae, plant cells, insect cells and animal cells. Expression vectors include, but are not limited to, cloning vectors, modified cloning vectors, specifically designed plasmids or viruses. Specifically designed vectors allow the shuttling of DNA between hosts, such as bacteria-yeast or bacteria-animal cells.

30 DNA encoding antibodies may also be cloned into an expression vector for expression in a host cell. Host cells may be prokaryotic or eukaryotic, including but not limited to bacteria, yeast, mammalian and insect cells and cell lines.

The expression vector may be introduced into host cells via any one of a number of techniques including but not limited to

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transformation, transfection, protoplast fusion, and electroporation.

Expression of cloned DNA may also be performed using *in vitro* produced synthetic mRNA. Synthetic mRNA can be efficiently translated in various cell-free systems, including but not limited to wheat germ extracts and reticulocyte extracts, as well as efficiently translated in cell based systems, including but not limited to microinjection into frog oocytes, with micro-injection into frog oocytes being preferred.

It is also well-known that there is a substantial amount of redundancy in the various codons which code for specific amino acids. Therefore, this invention is also directed to those DNA sequences which contain alternative codons which code for the eventual translation of the identical amino acid. For purposes of this specification, a sequence bearing one or more replaced codons will be defined as a degenerate variant.

The following examples are provided to further define the invention without, however, limiting the invention to the particulars of these examples.

20

## EXAMPLE 1

### Construction of mutations

Plasmid pP5Q was the starting vector for all mutagenic studies. Plasmid pP5Q is a derivative of p5H7 (Cambridge Antibodies). Plasmid pP5Q contains the VH and VL regions originally derived from MAbs 447 (Gormey *et al.*) cloned as a single chain fragment variable (scFv).

Table 1 lists some of the oligonucleotide primers used for site-directed mutagenesis of complementary determining region 3 (CDR3) of MAbs 447. Primers were synthesized on either a model 381A DNA Synthesizer (Applied Biosystems, Foster City, CA) or a Cyclone™ Plus DNA Synthesizer (MilliGen/Bioscience, Marlborough, MA). Mutagenesis was performed with the Transformer™ Mutagenesis Kit (CLONTECH, Palo Alto, CA)

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according to the manufacturer's instructions. All mutations were verified by DNA sequencing using the Sequenase® V2.0 DNA Sequencing Kit (United States Biochemical, Cleveland, OH).

5

Table 1

Primers:

Randomization of position 107:

10

CTC GGA GAC TCC C/GNN AAT CAT AAT AAA

Randomization of position 111:

GTA GTA GTA GTC C/GNN GGA GAC TCC CCG

15

Randomization of position 112:

GTC GTT GTA GTA GTA GTA C/GNN CTC GGA GAC

EXAMPLE 2

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Preparation of extracts and BIACore analysis of scFv Extracts:

25

Mutagenized plasmids were introduced by electroporation into bacterial strain *Escherichia coli* TG1 for expression. Single colonies were inoculated into 10 ml of 2X-YT (which contains per liter of water 16 g tryptone, 10 g yeast extract and 5 g sodium chloride) supplemented with 2% glucose. Cells were grown overnight at 30°C with vigorous shaking, collected by centrifugation in a Beckman GPR centrifuge at 2500 rpm, and resuspended in 10 ml of fresh 2X-YT supplemented with 1 mM isopropyl-beta-D-thiogalactopyranoside (IPTG) to induce expression. Cells were incubated at 30°C for an additional 5–6 hours with vigorous shaking, collected by centrifugation, resuspended in 1 ml of phosphate buffered saline: ethylenediametetraacetic acid (PBS:EDTA; 10 mM sodium phosphate pH7.0, 150 mM sodium chloride 1 mM EDTA), and incubated on ice for 30 minutes to

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release periplasmic proteins. Extracts were clarified by centrifugation and stored at 4°C until use.

EXAMPLE 3

5

Off-rate determinations of the scFv antibodies were determined using the BIACore system (Pharmacia Biosenser). HIV gp120 V3 loop peptides, Al-1 variant (Ala-1 peptide) were covalently immobilized on a carboxylated dextran/gold matrix via the primary amino group. The carboxyl-dextran matrix was first activated with N-ethyl-N'-(3-diethylaminopropyl)carbodiimide (EDC) and reacted with N-hydroxysuccinimide (NHS). HIV gp120 V3 loop peptides such as Ala-1 peptide were covalently immobilized via the free thiol of a cysteine placed at the N-terminus. These peptides were reacted with the EDC-NHS activated matrix which had been reacted with 2-(2-pyridinyldithio)ethaneamine. Remaining unreacted NHS-ester groups were displaced by addition of ethanolamine. EDTA extracts were added in a flow passing over the immobilized antigen. The refractive index changes, in the form of the surface plasmon resonance caused by the binding and subsequent dissociation of the scFv, were monitored continuously. Off-rates were calculated from the automatically collected data using the Pharmacia Kinetics Evaluation software.

25

EXAMPLE 4

Alanine scanning of CDR3 identifies residues which modulate scFv-antigen binding

30

Alanine scanning mutagenesis was used to identify residues within the VH CDR3 region of scFv clone P5Q critical for binding. It was hypothesized that effects on binding by alanine substitution would lead to four broad classes of effect: class i) slower off-rate; class ii) faster off-rate; class iii) loss of binding; and class iv) minor or no change in off-rate. Class i) and ii) were

- 10 -

operationally defined as critical. Class iii) was defined as obligatory. Class iv) was defined as noncritical.

The 27 positions that comprise VH CDR3 of scFv clone P5Q were individually changed to alanine by site-directed mutagenesis. Periplasmic extracts were prepared from the alanine replacement clones and assayed for off-rate determinations against the AL-1 gp120 V3 loop peptide (Fig. 1). Alanine substitutions at positions 107 and 111 resulted in 1.7 and 2.7 fold improvements in off-rate, respectively. These positions (class i) were judged critical and subsequently randomized to identify optimal residues. Alanine substitutions at positions 102, 112, 113, 114, and 118 led to faster off-rates (class ii); two of these positions were selected for further evaluation. Alanine substitution at positions 98, 101, 115, 116, 117, and 121 resulted in no binding (class iii). Alanine substitution at the remaining fourteen positions had only a minor effect on the off-rate (class iv). The class iii and iv positions were not evaluated further.

#### EXAMPLE 5

Randomization at critical positions to identify optimal amino acid solutions

---

The two critical class i) positions (107 and 111) were individually randomized to all amino acids, and off-rates against the AL-1 peptide determined. In addition, two class ii) positions (112 and 118) were also selected for randomization studies.

The results for position 107 are shown in Fig. 2. The slowest off-rate was observed with the negatively-charged glutamic acid, which decreased dissociation 2.5-fold. Substitution of other polar and charged amino acids had no significant effect on dissociation. With the exception of alanine, substitution with hydrophobic amino acid resulted in complete loss of binding. These results are consistent with the preponderance of surface ligand-contact residues being hydrophilic.

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Randomization of position 111 (Fig. 3) showed that the aromatic residues tyrosine and tryptophan produced the slowest off-rates (dissociation rates decreased 4.2 and 4.7-fold, respectively). However, substitution with any hydrophobic amino acids increased affinity relative to wild-type clone P5Q.

5 Class ii) positions 112 and 118 (faster off-rate upon alanine substitution) were also selected for amino acid randomization. For both position 112 (Fig. 4) and 118, the residues present in the original scFv P5Q, aspartic acid and asparagine, were 10 the best solutions.

#### EXAMPLE 6

##### Improvements at positions 107 and 111 are additive

15 A double mutant that combined the optimized residues at positions 107 (E) and 111 (W) was constructed to determine whether or not the individual improvements are additive. Figure 5 shows that the double mutant has an off-rate 9-fold slower than wild-type clone P5Q. The off-rate value approximates the product of the fold 20 improvements observed with the individual optimized residues (2.5 for 107E and 4.7 for 111W). One interpretation of this result is that for these two positions, the contributions to scFv-antigen affinity are independent and additive.

#### EXAMPLE 7

##### Method of making modified antibodies

An antibody is mutagenized by alanine scanning 30 mutagenesis to produce a modified antibody. The binding of the modified antibody to its antigen is determined. Binding determinations may be made by conventional methods and include off-rate measurements. Modified antibodies having desired characteristics are selected and maintained.

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EXAMPLE 8

Method of using modified antibodies

5       The modified antibodies or pharmaceutical compositions thereof are used for the prophylactic or therapeutic treatment of diseases caused by their antigen. Methods of treatment include, but are not limited to, intravenous or intraperitoneal injection of the modified antibody.

10

EXAMPLE 9

Diagnostic kit employing modified antibodies

15       The modified antibodies of Example 7 are used as reagents in diagnostic kits. The modified antibody reagents may be further modified through techniques which are well-known in the art, such as radiolabeling or enzyme-labeling. The diagnostic kit may be used to detect or characterize the antigens.

20

EXAMPLE 10

DNA encoding modified antibodies

25       The DNA encoding the modified antibody of Example 7 is used as a reagent for the production of modified antibodies. The DNA may be incorporated into an expression vector. The expression vector may be used to transform a host cell. Cultivation of the host cell under conditions suitable for the expression results in the production of modified antibody.

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EXAMPLE 11

DNA encoding modified antibodies

5       The DNA encoding the modified antibody of Example 7  
is used to detect DNA encoding the antigen in test samples. Methods  
of detection include, but are not limited to, hybridization under  
selective conditions. Test samples include, but are not limited to,  
samples of blood, cells, and tissues.

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EXAMPLE 12

Preparation of modified light chain immunoglobulins

15       The light chain of an immunoglobulin is mutagenized by  
alanine scanning mutagenesis to produce a modified immunoglobulin  
having modified binding characteristics. The modified immuno-  
globulin is used as a reagent for diagnostic kits or as a therapeutic  
agent.

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANT: LEWIS, CRAIG M.  
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(ii) TITLE OF INVENTION: IN VITRO ANTIBODY MATURATION

(iii) NUMBER OF SEQUENCES: 2

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(D) STATE: NJ  
(E) COUNTRY: USA  
(F) ZIP: 07065

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk  
(B) COMPUTER: IBM PC compatible  
(C) OPERATING SYSTEM: PC-DOS/MS-DOS  
(D) SOFTWARE: PatentIn Release #1.0, Version #1.25

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(C) CLASSIFICATION:

(viii) ATTORNEY/AGENT INFORMATION:

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(C) REFERENCE/DOCKET NUMBER: 19190P

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(B) TELEFAX: (908) 594-4720

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 816 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

GCCATGGCCG AGGTGCAGCT GGTGGAGTCT GGGGGAGGCT TGGTAAAGCC TGGGGGGTCC	60
CTCAGACTCA CCTGTGTAGC CTCTGGCTTC ACGTTCAGTG ATGTCTGGCT GAACTGGTC	120
CGCCAGGCC CAGGGAAGGG GCTGGAGTGG GTGGCCGTA TTAAAAGCGC CACTGATGGT	180
GGGACAACAG ACTACGCTGC ATCCGTGCAA GGCAGATTCA CCATCTCAAG AGATGACTCA	240
AAAAACACGC TATATCTGCA AATGAATAGC CTGAAAACCG AGGACACAGC CGTTTATTCC	300
TGCAAACACAG ATGGTTTTAT TATGATTCGG GGAGTCTCCG AGGACTACTA CTACTACTAC	360
AACGACGTTT GGGGCAAAGG GACCACGGTC ACCGTCTCCT CAGGTGCAGG CGGTTCAAGC	420
GGAGGTGGCT CTGGCGGTGG CGGATCGCAG TCTGTGTTGA CGCAGCCGCC CTCAGTGTCT	480
GC GGCCCCAG GACAGAACGGT CACCATCTCC TGCTCTGGAA GCAGCTCCAA CATTGGGAAT	540
AATTATGTAT TGTGGTACCA GCAGTTCCCA GGAACAGCCC CCAAACCTCCT CATTATGGC	600
AATAATAAGC GACCCCTCAGG GATTCCCTGAC CGATTCTCTG GCTCCAAGTC TGGCACGTCA	660
GCCACCCCTGG GCATCACCGG ACTCCAGACT GGGGACGAGG CCGATTATTT CTGCCCAACA	720
TGGGATAGCG GCCTGAGTGC TGATTGGGTG TTCGGCGGAG GGACCAAGCT GACCGTCCTA	780
GGTGCAGCCG CAGAACAAAA ACTCATCTCA GAAGAG	816

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 272 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Ala Met Ala Glu Val Glx Leu Val Glu Ser Gly Gly Gly Leu Val Lys			
1	5	10	15
Pro Gly Gly Ser Leu Arg Leu Thr Cys Val Ala Ser Gly Phe Thr Phe			
20	25	30	
Ser Asp Val Trp Leu Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu			
35	40	45	
Glu Trp Val Gly Arg Ile Lys Ser Ala Thr Asp Gly Gly Thr Thr Asp			
50	55	60	

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Tyr Ala Ala Ser Val Gln Gly Arg Phe Thr Ile Ser Arg Asp Asp Ser  
65 70 75 80

Lys Asn Thr Leu Tyr Leu Glx Met Asn Ser Leu Lys Thr Glu Asp Thr  
85 90 95

Ala Val Tyr Ser Cys Asn Thr Asp Gly Phe Ile Met Ile Arg Gly Val  
100 105 110

Ser Glu Asp Tyr Tyr Tyr Tyr Asn Asp Val Trp Gly Lys Gly Thr  
115 120 125

Thr Val Thr Ala Ser Ser Gly Ala Gly Gly Ser Gly Gly Gly Ser  
130 135 140

Gly Gly Gly Ser Gln Ser Val Leu Thr Gln Pro Pro Ser Val Ser Ala  
145 150 155 160

Ala Pro Gly Gln Lys Val Thr Ile Ser Cys Ser Gly Ser Ser Asn  
165 170 175

Ile Gly Asn Asn Tyr Val Leu Trp Tyr Gln Gln Phe Pro Gly Thr Ala  
180 185 190

Pro Lys Leu Leu Ile Tyr Gly Asn Asn Lys Arg Pro Ser Gly Ile Pro  
195 200 205

Asp Arg Phe Ser Gly Ser Lys Leu Leu Ile Tyr Gly Ala Thr Leu Gly  
210 215 220

Ile Thr Gly Leu Gln Thr Gly Asp Gln Ala Asp Tyr Phe Cys Ala Thr  
225 230 235 240

Trp Asp Ser Gly Leu Ser Ala Asp Trp Val Phe Gly Gly Thr Lys  
245 250 255

Leu Thr Val Leu Gly Ala Ala Ala Glu Gln Lys Leu Ile Ser Glu Glu  
260 265 270

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WHAT IS CLAIMED IS:

1. A DNA molecule encoding a modified antibody,  
the modified antibody being derived from a native antibody by  
alanine scanning mutagenesis and the modified antibody having  
binding characteristics different than binding characteristics of the  
native antibody.

2. The DNA molecule of Claim 1 wherein the native  
antibody is MAb447.

3. The DNA molecule of Claim 2, the DNA  
molecule being selected from the group consisting of P5Q, DNA  
encoding modified antibodies of Figures 1, 2, 3, 4, 5, combinations  
thereof, derivatives thereof and degenerate variants thereof.

4. A method of modifying an antibody to make an  
modified antibody comprising replacing at least one amino acid of  
the antibody with alanine to produce a modified antibody.

5. The method of Claim 4 wherein the modified  
antibody has improved binding characteristics.

6. Modified antibodies produced by the method of  
Claim 4 or homologues thereof.

7. The method of Claim 4 wherein the antibody is  
MAb447.

8. The method of Claim 7 wherein the amino acid  
replaced with alanine is located in complementary determining  
region 1, complementary determining region 2 or complementary  
determining region 3 of MAb447.

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9. The modified antibodies of Claim 6 selected from the group consisting of P5Q, the antibodies of Figures 1, 2, 3, 4, 5, combinations thereof, derivatives thereof, and homologues thereof.

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10. Diagnostic kits comprising the modified antibodies produced by the method of Claim 6.

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11. Diagnostic kits comprising the DNA molecules of Claim 1.

10

12. A pharmaceutical composition comprising at least one modified antibody of Claim 6 or DNA encoding at least one modified antibody of Claim 6 or combinations thereof.

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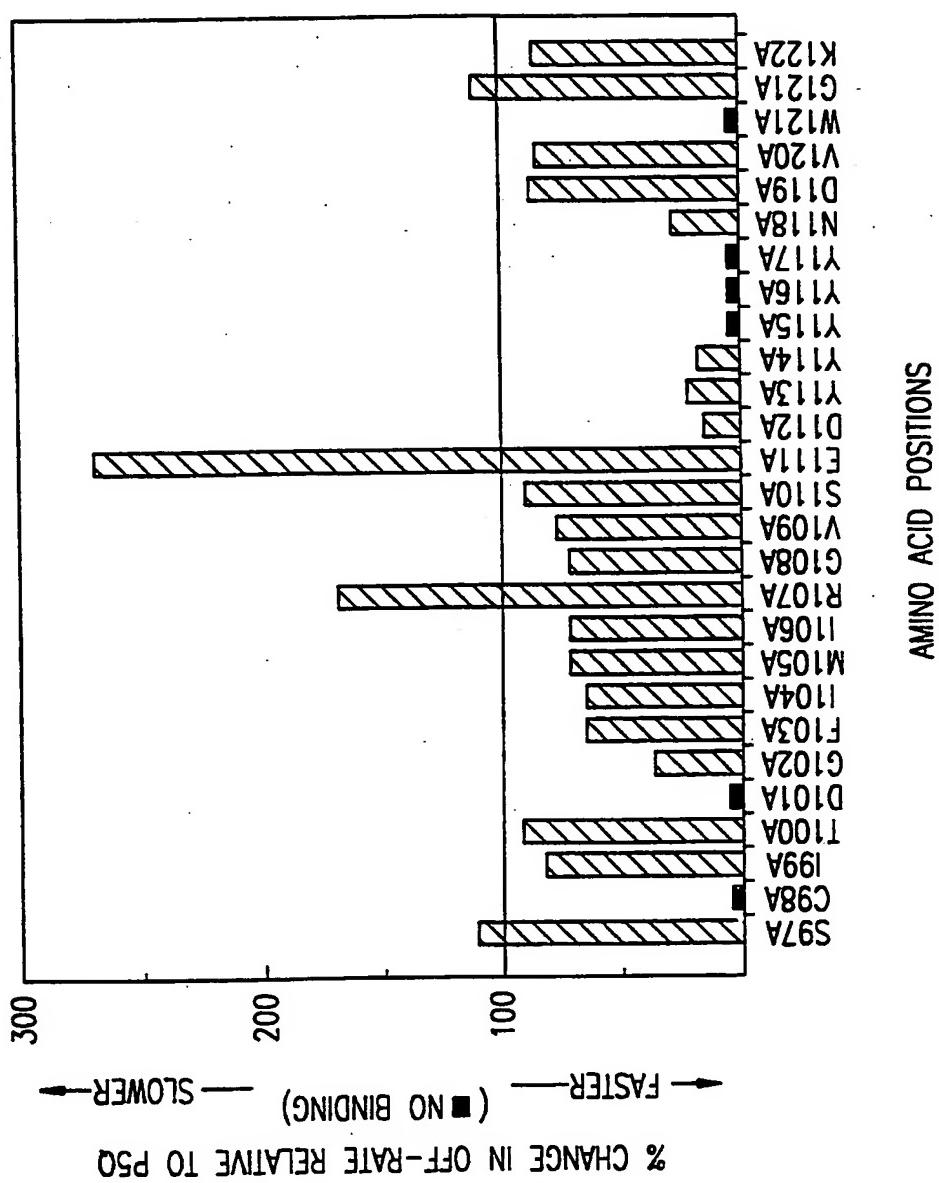


FIG. 1

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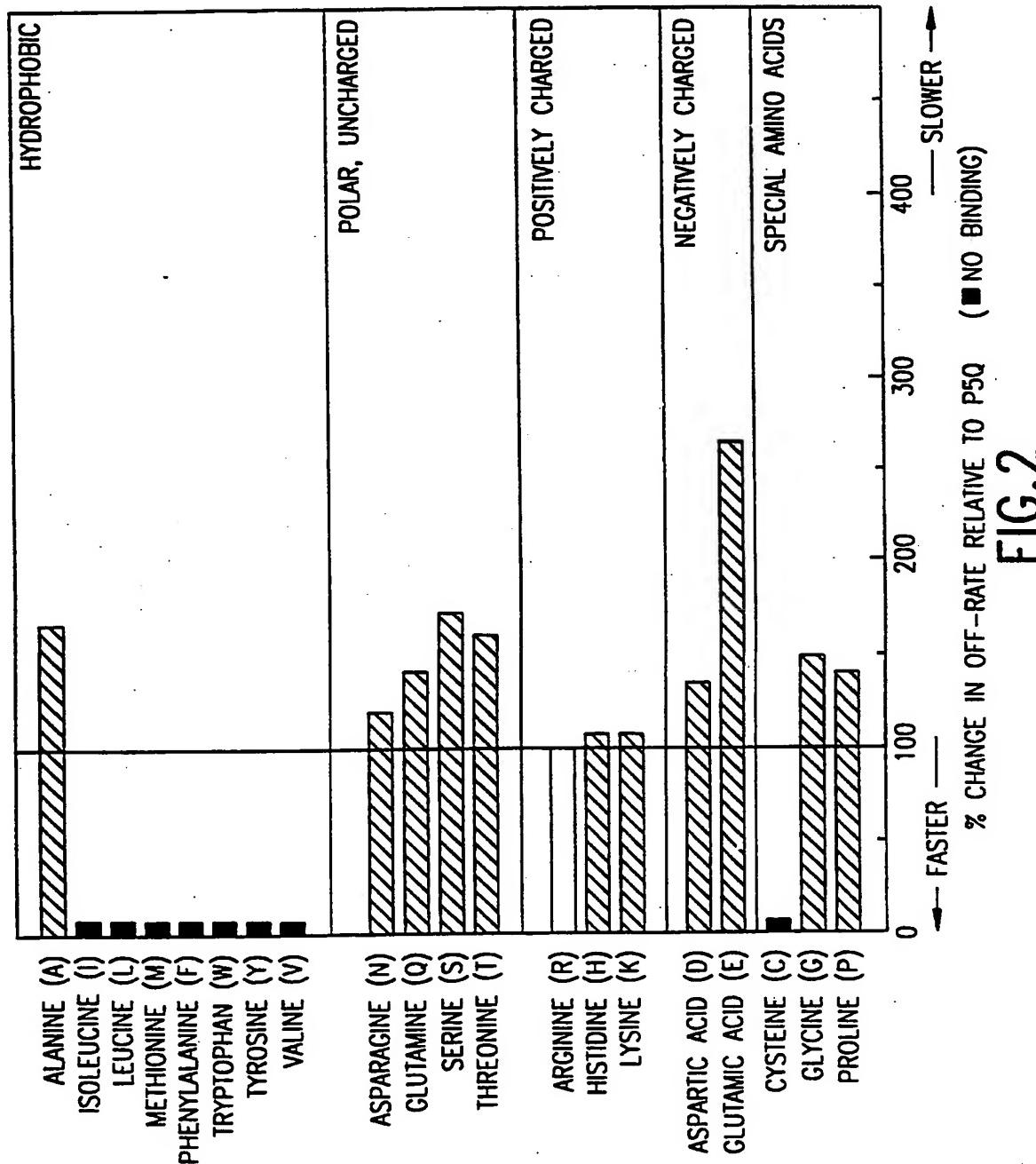


FIG. 2

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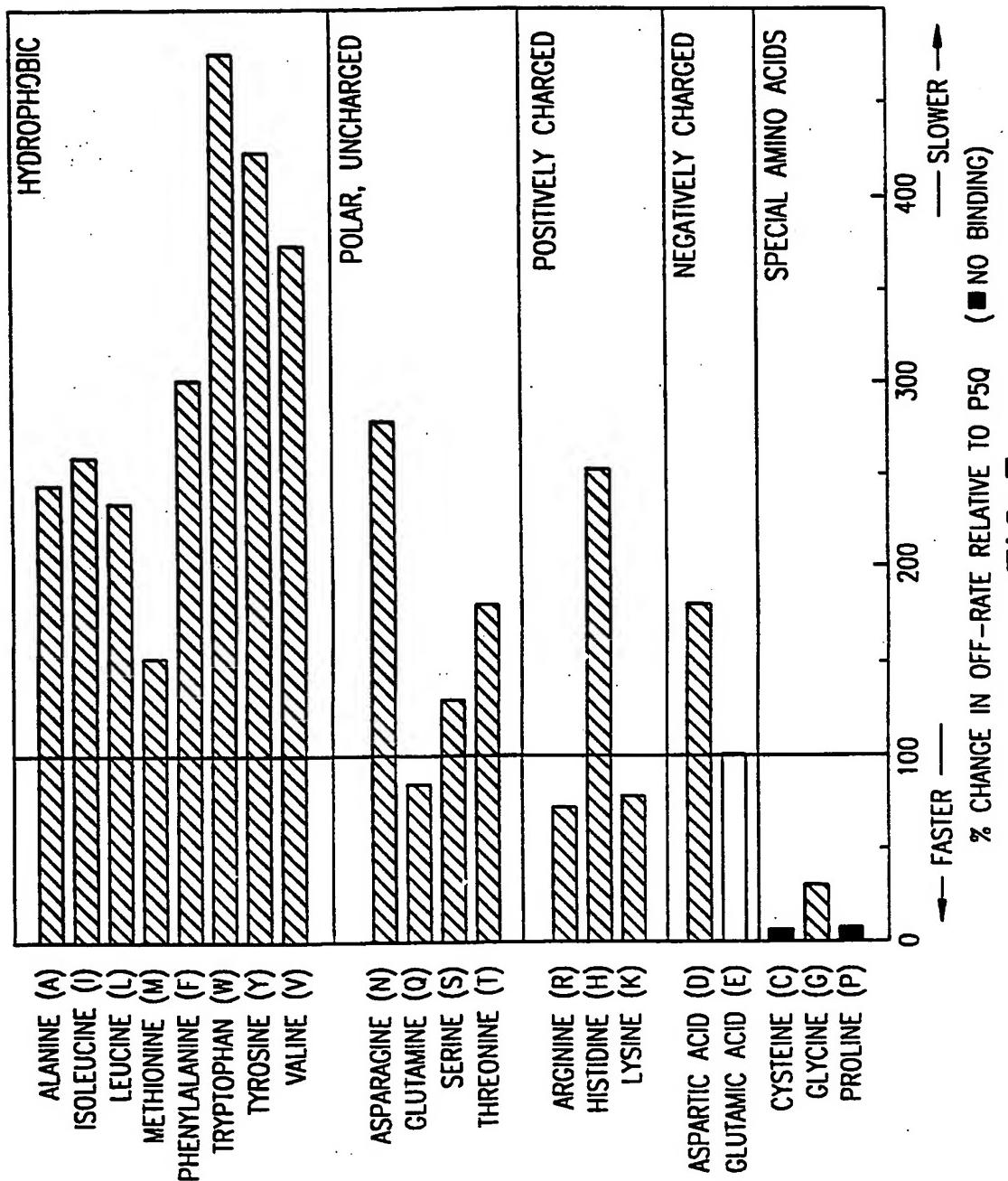


FIG. 3

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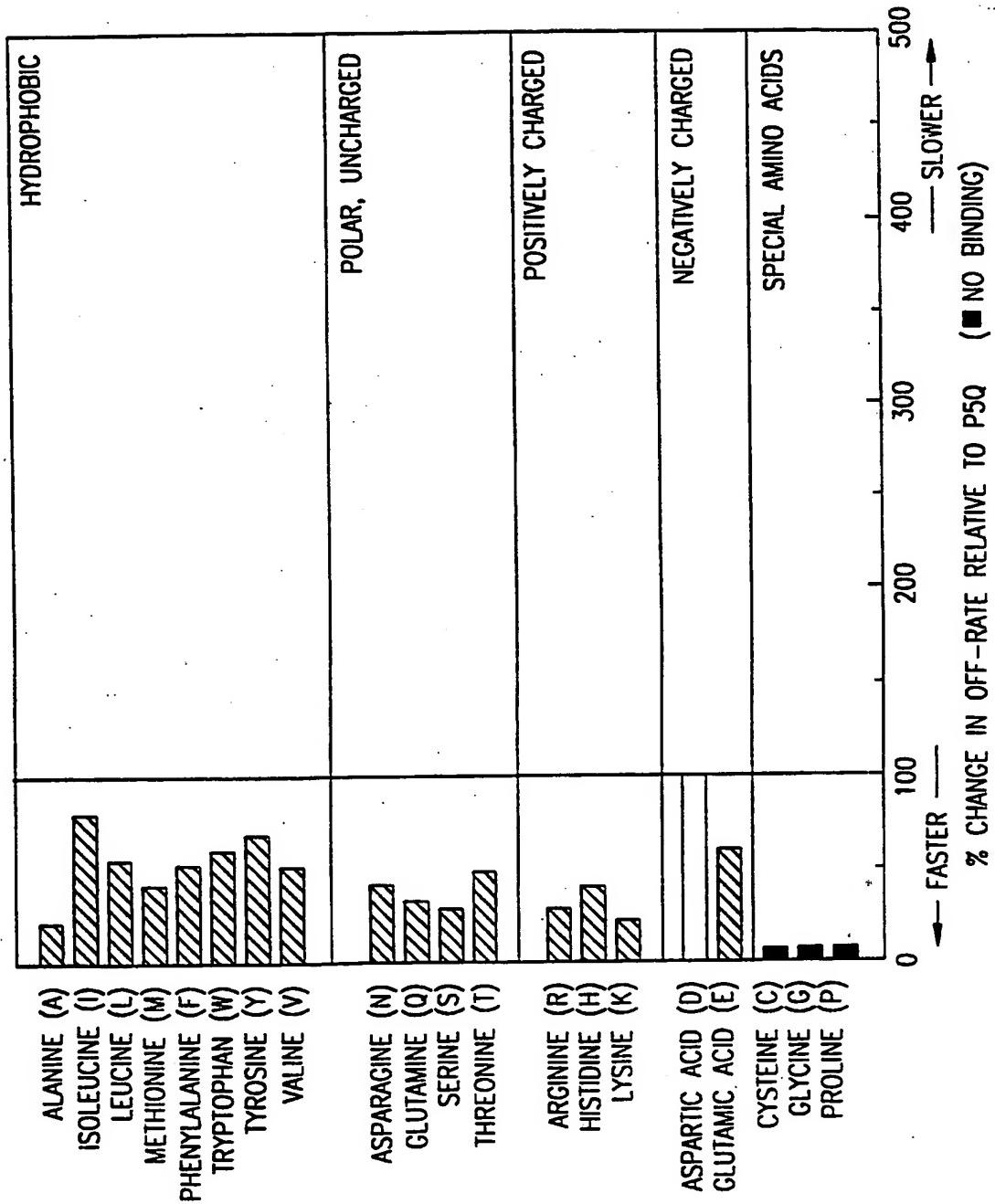


FIG. 4

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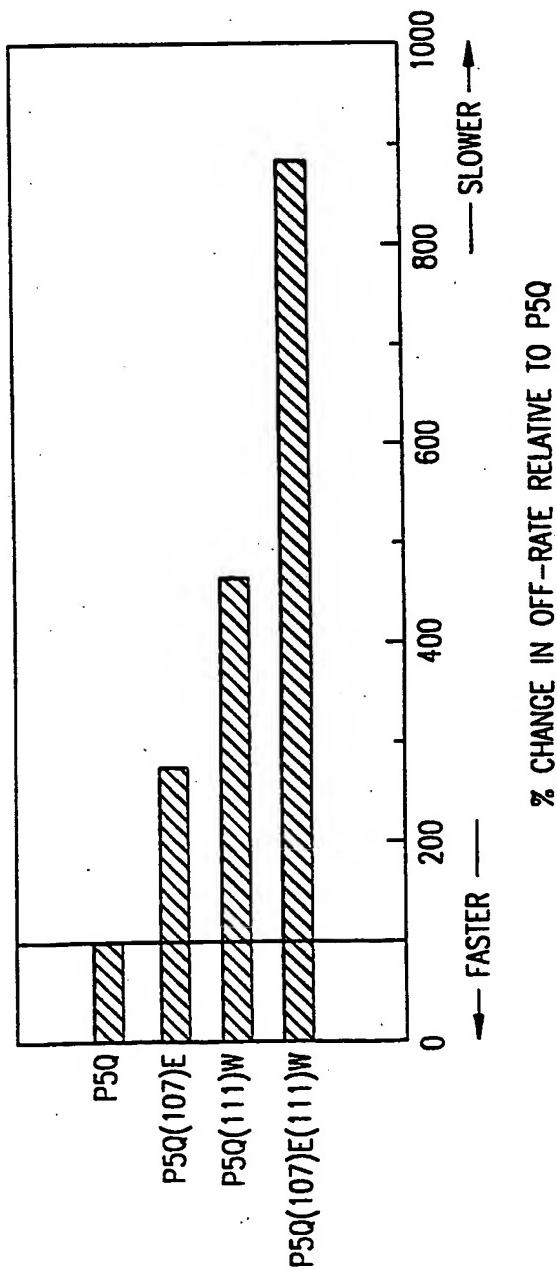


FIG. 5

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10	20	30	40	50	60
*	*	*	*	*	*
CCC ATG CCC GAC GTG CAG CTC CTC GAG TCT CCC CGA CGC TTG CTA AAG CCT CCC CGG TCC					
Ala Met Ala Glu Val Gln Leu Val Ser Gly Glu Gly Leu Val Lys Pro Gly Gly Ser					
70	80	90	100	110	120
*	*	*	*	*	*
CTC AGA CTC ACC TGT GTA GCC TCT CGC TTC ACG TTC AGT GAT GTC TGG CTG AAC TGG GTC					
Leu Arg Leu Thr Cys Val Ala Ser Gly Phe Thr Phe Ser Asp Val Trp Leu Asn Trp Val					
130	140	150	160	170	180
*	*	*	*	*	*
CCC CAG CCA CCC AAC CGG CTC GAG TGG GTC CGC CGT ATT AAA AGC GCC ACT GAT CGT					
Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Gly Arg Ile Lys Ser Ala Thr Asp Gly					
190	200	210	220	230	240
*	*	*	*	*	*
GGG ACA ACA GAC TAC GCT GCA TCC GTG CAA GCC AGA TTC ACC ATC TCA AGA GAT CAC TCA					
Gly Thr Thr Asp Tyr Ala Ala Ser Val Gln Gly Arg Phe Thr Ile Ser Arg Asp Asp Ser					

FIG.6d

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250	*	260	*	270	*	280	*	290	*	300	*
AAA	AAC	ACG	CTA	TAT	CTG	CAA	ATG	AAT	AGC	CTG	AAA
Lys	Asn	Thr	Leu	Tyr	Leu	Gln	Met	Asn	Ser	Leu	Lys
310		320	*	330	*	340	*	350	*	360	*
TGC	AAC	ACA	GAT	GGT	TTT	ATT	ATG	ATT	CGG	GGA	GTC
Cys	Asn	Thr	Asp	Gly	Phe	Ile	Met	Ile	Arg	Gly	Val
370		380	*	390	*	400	*	410	*	420	*
AAC	GAC	GTT	TGG	GGC	AAA	GGG	ACC	ACC	GTC	ACC	GTC
Asn	Asp	Val	Trp	Gly	Lys	Gly	Thr	Thr	Val	Thr	Ala
430		440	*	450	*	460	*	470	*	480	*
GCA	GGT	GGC	TCT	GGC	GGC	GCA	TCC	CAG	TCT	GTC	TCT
Gly	Gly	Gly	Ser	Gly	Gly	Gly	Ser	Gln	Ser	Val	Leu

FIG. 6b

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490	500	*	510	*	520	*	530	*	540	*										
CGG	GCC	CCA	CGA	CAG	AAG	GTC	ACC	ATC	TCC	TGC	TCT	GGG	AGC	ACC	TCC	AAC	ATT	GGG	AAT	
Ala	Ala	Pro	Gly	Gln	Lys	Val	Thr	Ile	Ser	Cys	Ser	Gly	Ser	Ser	Ser	Asn	Ile	Gly	Asn	
550	560	*	570	*	580	*	590	*	600	*										
AAT	TAT	GTA	TTG	TGG	TAC	CAG	CAG	TTC	CCA	GGG	ACA	GCC	CCC	AAA	CTC	CTC	ATT	TAT	GGC	
Asn	Tyr	Val	Leu	Trp	Tyr	Gln	Gln	Pro	Pro	Pro	Gly	Thr	Ala	Pro	Lys	Leu	Leu	Ile	Tyr	Gly
610	620	*	630	*	640	*	650	*	660	*										
AAT	AAT	AAG	CGA	CCC	TCA	GGG	ATT	CCT	GAC	CGA	TTC	TCT	GGC	TCC	AAG	TCT	GGC	ACG	TCA	
Asn	Asn	Lys	Arg	Pro	Ser	Gly	Ile	Pro	Asp	Arg	Phe	Ser	Gly	Ser	Lys	Ser	Gly	Thr	Ser	
670	680	*	690	*	700	*	710	*	720	*										
GGC	ACC	CTG	GGC	ATC	ACC	GGA	CTC	CAG	ACT	GGG	GAC	GAG	GGC	GAT	TAT	TTC	TGC	GCA	ACA	
Ala	Thr	Leu	Gly	Ile	Thr	Gly	Leu	Gln	Thr	Gly	Ser	Glu	Ala	Asp	Tyr	Phe	Cys	Ala	Thr	

FIG.6c

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730            740            750            760            770            780  
\*            \*            \*            \*            \*            \*  
TGG GAT ACC GCC CTC ACT GCT GAT TGG GTC TTC CCC GGA GGC ACC AAC CTC ACC GTC CTA  
Trp Asp Ser Gly Leu Ser Ala Asp Trp Val Phe Gly Gly Thr Lys Leu Thr Val Leu  
  
790            800            810            \*  
\*            \*            \*            \*  
GCT GCG CCC GCA GAA CAA AAA CTC ATC TCA GAA GAG  
Gly Ala Ala Glu Gln Lys Leu Ile Ser Glu Glu

FIG. 6d

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US95/02492

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(6) :C07K 16/00, 16/46; A61K 39/00; C12N 15/12, 15/13

US CL :424/133.1, 144.1; 536/23.53; 530/387.3

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 424/133.1, 144.1; 536/23.53; 530/387.3

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

SEQUENCE SEARCH, MEDLINE, EMBASE, LIFESCI, BIOSYS, WPI

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	J. IMMUNOLOGY, VOL. 150, NO. 2, ISSUED 15 JANUARY 1993, M.K. GORNY ET AL., "REPERTOIRE OF NEUTRALIZING HUMAN MONOClonAL ANTIBODIES SPECIFIC FOR THE V3 DOMAIN OF HIV-1 GP120", PAGES 635-643, SEE ENTIRE DOCUMENT.	1-12
Y	PROC. NATL. ACAD. SCI. USA, VOL. 87, ISSUED SEPTEMBER 1990, A. ASHKENAZI ET AL., "MAPPING OF THE CD4 BINDING SITE FOR HUMAN IMMUNODEFICIENCY VIRUS BY ALANINE-SCANNING MUTAGENESIS", PAGES 7150-7154, SEE ENTIRE DOCUMENT.	1-12
Y	SCIENCE, VOL. 244, ISSUED 02 JUNE 1989, B.C. CUNNINGHAM ET AL., "HIGH-RESOLUTION EPITOPE MAPPING OF hGH-RECEPTOR INTERACTIONS BY ALANINE-SCANNING MUTAGENESIS", PAGES 1081-1085, SEE ENTIRE DOCUMENT.	1-12

<input type="checkbox"/>	Further documents are listed in the continuation of Box C.	<input type="checkbox"/>	See patent family annex.
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*A*	document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E*	earlier document published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
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*O*	document referring to an oral disclosure, use, exhibition or other means		
*P*	document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search	Date of mailing of the international search report
06 MAY 1995	23 MAY 1995
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231	Authorized officer, <i>Christopher Eisenchenk</i> CHRISTOPHER EISENSCHENK
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